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(11) Publication number : **0 568 265 A2**

(12)

## EUROPEAN PATENT APPLICATION

(21) Application number : 93303126.2

(51) Int. Cl.<sup>5</sup> : **A61M 1/36, A61M 5/165**

(22) Date of filing : 21.04.93

(30) Priority : 30.04.92 US 876041

(43) Date of publication of application :  
03.11.93 Bulletin 93/44

(84) Designated Contracting States :  
AT BE CH DE DK ES FR GB IT LI NL PT SE

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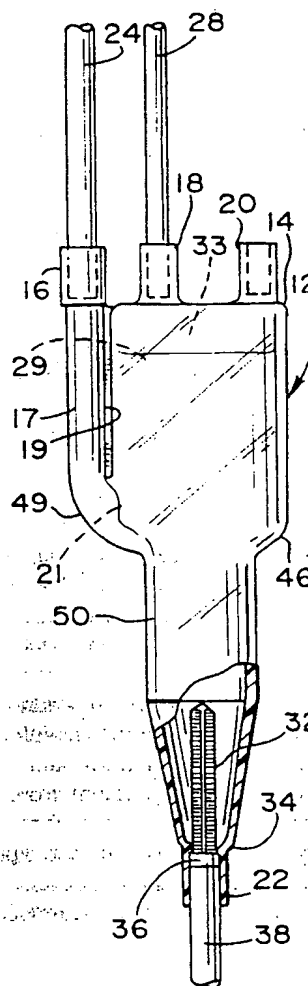
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(54) **Blow molded venous drip chamber for hemodialysis.**

(57) A blow molded, rigid plastic venous chamber (12) for hemodialysis, for use in a venous hemodialysis set (10). The chamber defines a plurality of first access ports (16,18,20) adjacent to a first end (14) thereof. A second access port (22) is provided at an opposed end. A tubular plastic filter (32) is positioned in close-fitting relation within the second access port to project into the chamber. Unique advantages are provided by such a blow-molded, filter-carrying chamber.

**Fig. 3**



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## BACKGROUND OF THE INVENTION

In the well-known medical procedure of hemodialysis, blood is collected from a vascular access of a patient, passing through a well-known arterial hemodialysis set by means of a blood pump, to the hemodialyzer. The blood passes through hemodialyzer into a well-known venous hemodialysis set, by which the blood is conveyed back to the vascular access of the patient. Pediatric flow rates are as low as 20ml/min while adult flow rates are as much as 650ml/min. This entire set-up is known as an extracorporeal circuit.

The venous hemodialysis set has a well-known venous blood chamber, a so-called drip chamber that is typically 80-120mm long and 17-25mm outer diameter, designed to contain typically 7-35ml of blood as a flow-through reservoir and operated such that blood does not completely fill the drip chamber. The chamber top has a blood inlet tube connection port in axial orientation communicating through the top to a blood inlet downspout in the main cavity of the chamber. The downspout length may vary such that the downspout outlet is either above the blood level (i.e. in the airspace), or beneath the blood level. Blood exits the venous chamber at the bottom by an outlet communicating axially through the bottom to a blood outlet tube connection port. A filter is placed within the chamber just above the blood outlet.

The venous drip chamber serves a number of functions:

1. Air bubbles contained in the incoming blood flow are forced by their buoyancy and reduction in velocity to rise to the air space. Flow out of the chamber is thereby degassed.
2. Blood pressure can be measured via a pressure monitor tube communicating through the chamber top into the air space.
3. Air foam is also prevented from exiting the chamber by an ultrasonic air/foam detector attached to the outside of the drip chamber or the outside of the tubing just below the venous chamber. The detector stops the blood pump and causes a clamp to close below the venous drip chamber if air or foam is sensed.
4. Medicament tube access ports or injection sites are located on the chamber top, communicating with the air space.
5. Clots, etc. are prevented from leaving the chamber by the presence of a filter adjacent the exit of the chamber.

Currently, venous chambers are injection molded of flexible or rigid plastic, typically comprising a gently conical shape with an open end which is closed with a solvent-sealed top cap, after a plastic filter has been inserted through the open end and placed into snap-fit relation with the chamber near the bottom end. Snap-fit is required because, typically, the filter ma-

terial is not compatible for solvent bonding with the chamber material. Thus it is retained by detents on the inside of the chamber. This is an imperfect solution since:

1. Such a filter can occasionally come loose (when blood pressure expands the flexible chamber) and/or.
2. The space between the filter attachment ring and the chamber wall fills with blood which stagnates, promoting clots. The scientific literature (e.g. Ogden) reports such filters actually generate more clots than they trap.
3. Assembly is difficult and costly because the filter must be inserted with enough force to surmount the detents.

Disadvantages of injection molded chambers and top caps include:

1. Leaks between the top cap and the chamber body.
2. Clots promoted by blood foaming in its free-fall from a short downspout at flow-rates typically above 250ml/min. The common name of "drip chamber" came about because at low flow rates the blood "dripped" from a short downspout opening to the blood level. More recently, flow rates of up to 650ml/min have become possible because of newer hemodialyzers. But short downspouts foam severely at fast flow rates due to turbulence and cavitation caused by the rapid, continuous free fall of blood. Foam leads to increased clots due the greater surface area of an air/blood interface.
3. Clots promoted by blood stagnating in the space between long downspouts and the chamber wall. Long downspouts with an outlet beneath the blood level have been developed more recently in an attempt to reduce turbulence, in the same manner a hose dipped into a bucket of soapy water will not foam. However, blood stagnates in the tight space between the downspout and the chamber wall. Stagnation frequently leads to clots.
4. Clots promoted by blood stagnating above the outlet of a long downspout. At slower flow rates, long downspouts are counterproductive because blood stagnates in the blood layer above the downspout outlet, resulting typically in clotting.
5. Excessive air/foam alarms at high flowrates with long downspouts. Entrained air bubbles in the incoming blood flow is projected far enough down into the chamber such that they are sensed by the air/foam detector, creating many alarms and stopping the blood pump. It is well known that clotting increases whenever the blood flow is stopped.
6. Injection molding typically provides a structure having fairly sharp angles and edges, which is generally undesirable in blood handling apparatus.

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## DESCRIPTION OF THE INVENTION

Blowmolding a venous chamber overcomes many of these problems. Arterial chambers have long been blowmolded, such as Swan et al. U.S. Patent No. 4,666,598. These have many advantages over injection molded chambers:

1. Blowmolding creates a one-piece chamber, eliminating the prevalence of leaks from bonding two pieces together.

2. The blood inlet and outlet may be molded in a variety of shapes and angles, not limited to the axial orientation of injection molding. Also, the opening of the inlet (or outlet) into the chamber cavity may be in the chamber sidewall or chamber bottom rather than in a downspout separate from the chamber wall. Thus, no stagnant areas are created. Further, the inlet blood tube connecting port may be remote from the inlet opening to the chamber cavity. For example, the inlet blood tube connecting port may be in lateral relation to the chamber top while the inlet to the chamber cavity may be at the bottom, connected to the tube port via a conduit molded laterally to the chamber body.

It has long been known that a bottom inlet opening (causing upward flow into the chamber cavity) eliminates most causes of arterial chamber foaming, stagnation and clotting. The inlet flow is upward away from the bottom outlet (thus directing any entrained bubbles to the surface). No separate "upspout" within the chamber cavity is required, thus eliminating stagnation between the wall and chamber. And blood enters beneath the surface of the blood, eliminating any free-fall of blood, yet keeping all the blood in the chamber in gentle mixing eliminating stagnation.

3. Non-tubular shapes may be chosen to aid in preventing stagnant areas or turbulent areas. Many of these shapes are impossible by injection molding.

These advantages, however, have not been available before in a venous chamber:

1. Heretofore there has been no known way to place a filter into a blow molded chamber. Venous chambers require a filter.

2. Venous chambers must be of flexible material if they are to fit properly to air/foam detector probes. A flexible chamber is more difficult to blow mold as the parison sags more easily before it is blown.

3. A blood inlet opening at the bottom results in a non-tubular shape which cannot fit in current chamber holders for air/foam detectors.

By this invention, a blow-molded, flexible or semi-rigid plastic venous chamber for hemodialysis is

provided. The chamber defines a plurality of first access ports substantially adjacent to a first chamber and, one of which is a blood tube inlet port. A blood tube exit port is provided at substantially the opposed end of the chamber. By this invention, a tubular plastic filter is positioned in close-fitting relation within the exit port to project into the chamber.

By this means, the desired blow molded venous chamber may be provided, while carrying a filter in a manner which has not been previously successfully accomplished.

The blowmolded chamber may define an outwardly-facing annular step with the exit port. The tubular plastic filter defines an outer end portion that carries a radially outwardly projecting annular flange, proportioned to slidingly pass through the exit port, but to engage the annular step to limit inward travel of the filter to a predetermined, innermost position.

Additionally, flexible tubing may be sealed, typically solvent sealed, to the exit port and within the port at a position outwardly of the plastic filter. By this means, the plastic filter may be retained within the exit port by the presence of the tubing.

Typically, the blood inlet port may be positioned in separate, lateral relation to the chamber, with a communicating conduit extending from the blood inlet port at the first end of the chamber toward the opposed end in such separate, lateral relation. This blood inlet conduit enters the cavity of the chamber at a position below the first end, either in the sidewall or in the opposed, second end. By this means, a blow molded chamber may carry at least three access ports, while being readily manufacturable by flow molding technology, and without resulting in a blow molded chamber of unduly enlarged cross section.

Preferably, a portion of the blow molded venous chamber adjacent its first end is relatively flat and wide in cross section, when compared with a portion of the chamber below where the blood inlet conduit enters the cavity of the chamber. By this means a number of benefits accrue:

1. The flattened, upper portion facilitates the wide spacing of the first ports in the first end of the blowmolded chamber,

2. The tubular, lower portion allows the chamber to be placed in currently available air/foam detector equipment, which require a round, cross sectional fitment area.

3. The flattened, upper portion, in conjunction with a sidewall blood inlet to the chamber cavity, promotes gentle mixing of the blood at high or low flow rates.

Additionally, the blow molding technology which is utilized herein can provide blood-contacting inner portions thereof which are essentially free of sharp angles and edges. This facilitates the desirable, gentle handling of the blood as it passes through the hemodialysis set to which the chamber of this invention.

is attached.

The blow-molded flexible plastic venous chamber may be manufactured for use as part of a venous flow set for hemodialysis by blow molding a chamber having a plurality of first access ports adjacent one end; one of which is a blood tube connection port, and a second blood tube connection port at its other end. The blow molded chamber is then cooled, and a tubular plastic filter is inserted into the blood tube exit port in close-fitting relation, so that the filter projects into the blow molded chamber. The filter may be secured in the chamber.

Additionally, there may be formed in the blow molded chamber during the molding process an outwardly facing, annular step within the second access port. The tubular plastic filter may define an out end portion that carries a radially outwardly projecting annular flange which is proportioned to slidably pass through the exit port, but which is large enough to engage the annular step. This limits the inward travel of the filter to a predetermined, innermost position. Then, blood tubing of a set may be sealed in the exit port behind the filter and annular step, to secure the filter in position.

The plastic venous chamber of this invention may then be assembled into a venous set for hemodialysis, to exhibit the numerous advantages discussed.

In another aspect of this invention, venous hemodialysis blood lines typically have an injection site for the administration of drugs and the like, and for sampling. Traditional venous lines have injection sites mounted on:

- a. The chamber top cap and communication with the airspace above the blood level, or
- b. The sidewall of the chamber and communicating with the airspace above the blood level, as described in Heath et al., or
- c. The venous blood tubing upstream of the venous chamber.

Traditionally, any of these injection sites were suitable because they satisfied the nursing dictum that no injections be given below the air detector/line clamp assembly, in order to avoid the risk of air emboli given to the patient.

Today, however, important but expensive new drugs such as EPO (erythropoietin) cannot be administered interdialytically to one of these venous injection sites, because it is feared the drug could be trapped in the venous filter or chamber sidewalls where blood velocity is reduced. Nurses are now having to risk giving EPO and other drugs directly into the fistula needle, with the attendant risk to the patient of air emboli.

With a hemodialysis machine having an air detector/line clamp assembly mounted on the venous line typically at least two inches below a venous chamber, it is possible to design a venous bloodline whereby an injection site is placed in the tubing downstream of the

venous chamber, yet upstream of the air detector/line clamp.

By this means it is possible to inject valuable drugs interdialytically in a safe manner.

## DESCRIPTION OF THE DRAWINGS

In the drawings, Fig. 1 is a plan view of a venous hemodialysis set which carries a blow-molded chamber of this invention;

Fig. 2 is an enlarged, side elevational view of the chamber of Fig. 1, shown in the process of being blow molded;

Fig. 3 is an elevational view, rotated 90 degrees from the view of Fig. 2, of the chamber after blow molding, with the filter inserted, and also showing connection with the various tubings of the venous set;

Fig. 4 is a transverse sectional view taken along line 4-4 of Fig. 1;

Fig. 5 is an enlarged, elevational view of a blow molded chamber generally tubular in shape;

Fig. 6 is an enlarged, top end view of the blow molded chamber of Fig. 5;

Fig. 7 is an enlarged, elevational view of a blow molded chamber with the blood inlet conduit entering the chamber cavity from the second opposed end, with the filter inserted;

Fig. 8 is an enlarged, side elevational view of the chamber of Fig. 7;

Fig. 9 is an enlarged, top end view of the chamber of Fig. 7;

Fig. 10 is an enlarged, elevational view of a blow molded cassette carrying an arterial and venous chamber;

Fig. 11 is a side elevational view of the cassette of Fig. 10;

Fig. 12 is a top, plan view of the cassette of Fig. 10; and

Fig. 13 is a partially schematic view of a dialysis system showing another aspect of this invention.

## DESCRIPTION OF SPECIFIC EMBODIMENTS

Referring to the drawings, Fig. 1 shows a venous hemodialysis set 10 which may be generally conventional in design except for blow molded plastic chamber 12 of this invention and its relationship with set 10.

Blow molded plastic chamber 12, as shown, comprises a first end 14 defining three separate first access ports 16, 18, 20.

Plastic chamber 12 also defines a second access port 22 at the end opposed to the first end 14.

Blow molded venous chamber 12 may be blow molded from a parison of flexible or rigid, plastic such as rigid polyvinyl chloride, or poly(ethylene terephthalate), particularly a glycol-modified variety of the latter. Suitable plastic formulations for this purpose are

commercially available.

Access port 16 of chamber 12 communicates with flexible tubing 24, with tubing 24 typically being sealed within port 16. Tubing 24 at its other end carries a conventional connector 26, which is adapted for connection with the blood outlet port of a hemodialyzer.

Access port 18 is connected to pressure monitor tubing 28, which carries at its opposed end a conventional connector 30 for a pressure monitor.

Access port 20 may comprise a conventional needle pierceable partition of resealable elastomer, of a conventional design.

Second access port 22 is shown to carry a tubular plastic filter 32, which is proportioned to be inserted through second access port 22, as shown in Fig. 3. The interior of second access port 22 defines an internal, annular step 34. Rear flange 36 of filter 32 engages the outer surface of step 34, so that it is restrained from further inward motion into chamber 12. Flexible tubing 38 also enters into port 22 within the port at a position outwardly from plastic filter 32 and flange 36. Thus, the inner end of tubing 38 causes plastic filter 32 to be retained within second access port 22 by the presence of such tubing. Tubing 38 may be conventionally solvent sealed in its position in second port 22.

The opposed end of tubing 38 defines a conventional connector 40 for connection to a fistula needle, which, in turn, penetrates the fistula of the patient for return of dialyzed blood.

Injection site 42 is also carried on tubing 38, and may be of conventional design.

Access port 16 is shown to be carried on a separate, generally longitudinal arm 17 of chamber 12, which is separated from the remainder of chamber 12 by partition line 19. Partition line 19 is formed during the blow molding process so that tubular section 17 is approximately cylindrical in shape. Then, tubular section 17 communicates with the remainder of the chamber 12 at a lateral flow position 21 which is between the first end 14 and opposed end 22. By this means, a third port may be provided to the first end of chamber 12 without undue crowding of the respective ports.

Turning particularly to Fig. 2, the shape of the molded chamber 12 from the side is shown. It can be seen that the area adjacent first end 14 is relatively flat and wide in cross section, by comparison of Figs. 2 and 3, when compared with a portion 50 of the chamber which is spaced from first end 14, and typically which is adjacent opposed end 22. This facilitates the desired width and shape of portion 50 to fit into currently available machines with air/foam detectors requiring such a width and shape, while the area adjacent first end 14 does not provide undue volume or taking excessive amount of space to accommodate the three ports at first end 14.

Also, Fig. 2 shows the blow molding process of chamber 12 from a hot parison which is being repeatedly applied to the blow mold halves 42, 44 by a conventional extrusion means. As is conventional, the blow mold halves open, and the hot parison from the extruder is advanced into the space between blow mold halves 42, 44. The blow mold halves then close, pinching the parison at their respective ends 48, 51, or, additionally their sides 46, 49, while also forming partition 19 in the molding chamber. Then, compressed air or the like is conventionally inserted through blow tube 53 leading to first ports 16, 18, 20 and second port 22 respectively, to inflate the parison, so that the parison expands to the confines of the molding chamber defined with the blow mold halves 42, 44. This position is held until the parison has cooled sufficiently to form the desired chamber 12 of this invention.

Following trimming by conventional means, as shown in Fig. 3, plastic filter 32 may be inserted through second port 22, followed by insertion of tubing 38 which is sealed in place so that flange 36 is trapped between step 34 and tubing 38 for retention of the plastic filter in desired position.

The remaining tubings 24, 28 may be sealed into their respective ports, and the components of the injection site may be applied to port 20.

Thus, by this means, a venous hemodialysis set may be manufactured in which a blow molded chamber, having an inserted filter 32, is provided. The significant advantages of such a chamber are as discussed above. Specifically, the chamber of this invention may be made substantially free of sharp angles and edges at its blood-contacting inner portions. This is easily accomplished in blow molding as an inherent characteristic thereof, while such is not the case with respect to injection molded components. Also, the specific shape of the chamber of this invention permits chamber 12 to be of relatively small size and internal volume, while providing adequate cross section to accommodate three ports on first end 14. The entry of blood through port 16, 17 into a flattened, central portion of the chamber by smooth, radiused horizontal inlet 21 provides advantages, in that blood velocity is dissipated in its horizontal travel across the flattened cross section when it is turned upward by the gentle, radiused curve of shoulder 27. Flow will form a gentle counterclockwise pattern in the flattened section adjacent the first end. Any entrained air bubbles will outgass at the blood surface 29 to join the airspace 33. Further, at low flow rates the blood above inlet 21 will not become stagnant owing to the counterclockwise pattern, nor at high flow rates will undue turbulence be allowed as the velocity is dissipated. Further, there is no stagnation area between the conduit 17 and chamber wall as there is between an injection molded downspout and chamber wall. This invention allows one chamber suitable for high or

low flows, for all current machines.

Figs. 5 and 6 show a chamber similar to Figs. 1-4 except the cross sectional shape is substantially cylindrical in nature throughout its length.

Referring to Figs. 7 through 9, there is shown a blow molded venous chamber suitable for a class of dialysis machine where the air/foam detector is mounted below the chamber and its ultrasound probes are mounted on the tubing below the chamber. Here, blow molding may be of rigid plastic because the ultrasound probes do not touch the chamber wall, and the blood inlet to the main cavity of the chamber may be at the end opposed to the first end 114. The entry of blood through access port 118 is shown to be carried on a separate, generally longitudinal arm 110 of chamber 106, which is separated from the remainder of chamber 106 by partition line 120. Partition line 120 is formed during the blow molding process so that tubular section 110 is approximately cylindrical in shape. Then, tubular section 110 communicates with the remainder of the chamber 106 at a position opposite of the first end 114.

Second access port 132 is shown to carry a tubular plastic filter 138 which is proportioned to be inserted through second access port 132, and retained in a manner similar to filter 32 of Fig. 3. A partition 111 is also formed between blood inlet 129 and blood outlet 128. The partition 111 may be curved or pointed away from outlet 128. Looking at Fig. 8 in conjunction with Fig. 9, one can see that inlet 129 and outlet 128 are of preferably outwardly tapering cross section. In the case of blood entering the chamber via conduit 110, the velocity is dissipated by U-turn thereof as well as the outwardly tapering cross section in inlet 129.

Rigid material may be used for these reservoirs. Such material typically maintains its shape as a hot parison much better than flexible material. Savings in time and cost are provided. Also, by the inventive chamber shape, the blood flow is gentle, unturbulent, but efficiently degassed before exiting the chamber.

In Fig. 10 is shown a blow molded cassette incorporating an inventive venous chamber. This blow molded cassette comprises a single parison having a pair of spaced chambers, with chamber 204 being typically the arterial prepump chamber, and chamber 206 being typically the venous chamber. Blood inlet 230 receives blood from an arterial connector such as connector 26, while blood outlet 228 connects directly to pump tubing 229. Likewise for venous chamber 206, blood comes in from the venous connector of the dialyzer to inlet port 218, which connects to a separate conduit 210 that extends the length of the chambers 204, 206, forms a U-turn at 211, and enters the venous chamber 206. Blood outlet 232 from the venous chamber passes through filter 250 which may be of a structure and mounting similar to filter 32 in Figs. 2 and 3, with the blood traveling from there through

tubing 233 to the venous patient connector. Depending upon whether the system is used in a prepump or a postpump manner, saline or heparin can be inserted into inlets 216, 224, or 226. Conduit 208 which is separate from the chambers extends into the port area 230 of chamber 204. Flattened areas 220 are also defined by the blow mold to provide the discrete chambers 204, 206, and conduits 208, 210.

Thus, by the embodiment of Figs. 10-12 it can be seen that the prepump chamber and the postpump chamber as well as their various ports, can be formed into a single blowmolded cassette from a single parison.

Referring to Fig. 13, another invention of this application is disclosed. A somewhat schematic view of a dialyzer system is disclosed, with a conventional dialysis machine, specifically the Cobe Centry Two dialysis machine, carrying dialysis arterial and venous sets, which are of conventional design except as otherwise described herein.

An injection site 250 is positioned in the dialysis venous set downstream of venous chamber 252, but upstream of the position of the set tubing which passes through bubble detector 254 and downstream clamp 256, both of which are part of the hardware of the dialysis machine 258. By this modification, as described above, the administration of new drugs such as EPO can be effected with the advantages as described above.

The above has been offered for illustrative purposes only, and is not intended to limit the scope of the invention of this application, which is as defined in the claims below.

## Claims

1. A blow-molded plastic blood chamber for hemodialysis, said chamber defining a first access port adjacent a first end which is an inlet blood tube connecting port, an outlet blood tube connecting port at an opposed chamber end, and a tubular plastic filter positioned in close-fitting relation within said outlet port to project into said chamber.
2. The blow-molded chamber of Claim 1 in which an indentation is defined within said outlet port, said tubular plastic filter defining an outer end portion that carries a radially outwardly projecting, annular flange proportioned to slidably pass through said second access port, but to engage said indentation to limit inward travel of said filter to a predetermined, innermost position.
3. The blow-molded chamber of Claim 2 in which flexible tubing is sealed to said outlet port and within said outlet port at a position outwardly of

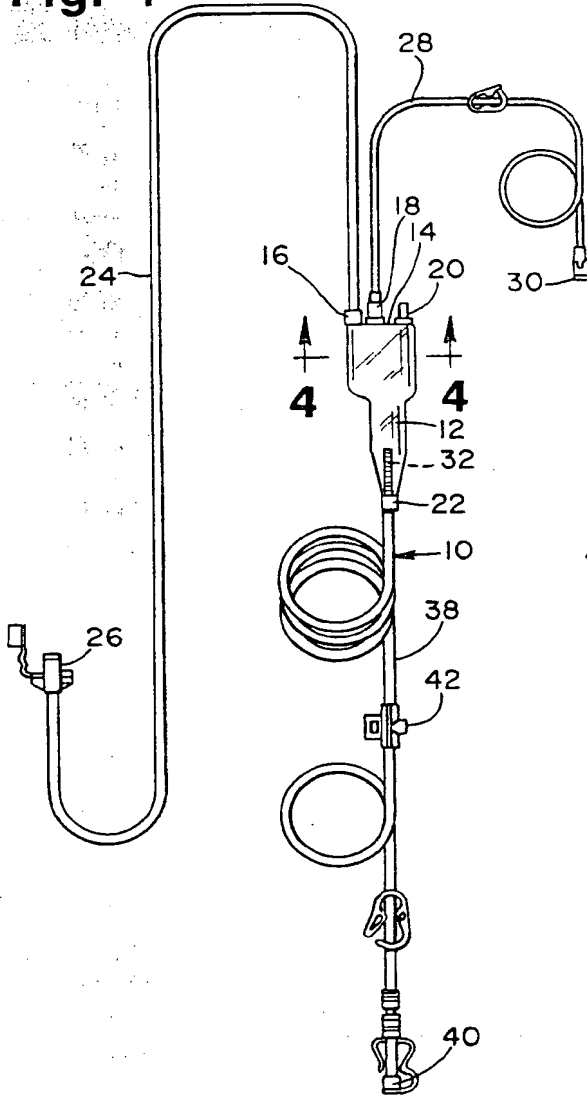
said plastic filter, whereby said flange and plastic filter is retained within said outlet port by the presence of said tubing.

4. The blow molded chamber of Claim 1 in which the inlet port is positioned in separate, lateral relation to said chamber, and from which extends a conduit toward said opposed end in said separate, lateral relation, while also communicating with said chamber at a position between said first and opposed ends.
5. The blow molded chamber of Claim 4 in which the conduit inlet communicates in radiussed horizontal direction.
6. The blow molded chamber of Claim 1 in which a portion of said chamber adjacent said first end is relatively oval in cross section, when compared with a portion of said chamber adjacent said opposed end.
7. The blow molded chamber of Claim 1 in which the inlet port is positioned in separate, lateral relation to said chamber and from which extends a conduit toward said opposed end in said separate, lateral relation, while also communicating with said chamber at said opposed end.
8. The blow molded chamber of Claim 7 in which the inlet port and outlet port are of the same height, by means of a partition positioned between them.
9. The blow molded chamber of Claim 1 in which three of said first access ports are present.
10. A venous hemodialysis set which includes at least one blow molded plastic venous chamber of Claim 1.
11. A blow-molded, rigid, plastic, venous chamber for hemodialysis, said chamber defining a plurality of first access ports adjacent a first end, a second access port at an opposed end, and a tubular, plastic filter positioned in close-fitting relation within said second access port to project into said chamber, at least one of said first access ports being positioned in separate, lateral relation to said chamber and extending from said first end toward said opposed end in separate, lateral relation, while also communicating with said chamber at a position between said first and opposed ends, and in which a portion of said chamber adjacent said first end is relatively flat and wide in cross section, when compared with a portion of said chamber adjacent said opposed end.
12. The blow-molded chamber of Claim 11 in which

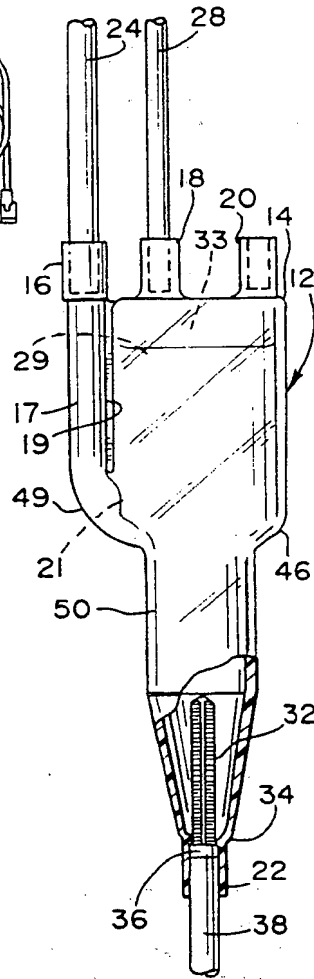
an outwardly-facing annular step is defined within said second access port, a tubular plastic filter defining an outer end portion that carries a radially outwardly projecting, annular flange proportioned to slidably pass through said second access port, but to engage said annular step to limit inward travel of said filter to a predetermined, innermost position.

13. The blow-molded chamber of Claim 12 in which flexible tubing is sealed to said second access port and within said port at a position outwardly of said plastic filter, whereby said plastic filter is retained within said second access port by the presence of said tubing.
14. A venous hemodialysis set which includes at least one blow-molded, rigid, plastic venous chamber of Claim 13.
15. The venous hemodialysis set of Claim 14 in which said blow molded chamber carries three of said first access ports.
16. The venous hemodialysis set of Claim 15 in which the blood contacting inner portions of said blow-molded chamber are essentially free of sharp angles and edges.
17. In a dialysis system which comprises dialysis set means mounted in a dialysis machine, said dialysis set means comprising a venous chamber, the improvement comprising an injection site positioned downstream of said venous chamber, but upstream of the bubble detector and downstream clamp of said dialysis apparatus, whereby drugs may be administered through the set to the patient downstream of the venous chamber.

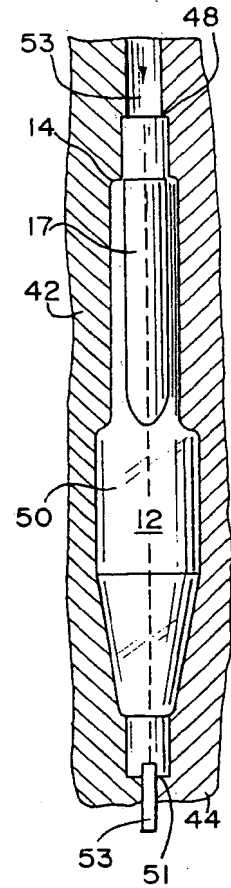
**Fig. 1**



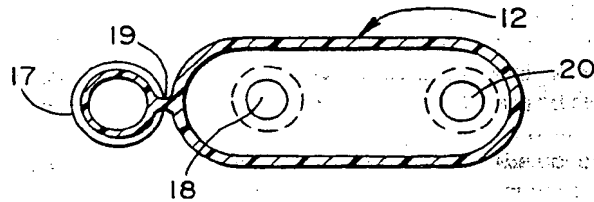
**Fig. 3**



**Fig. 2**

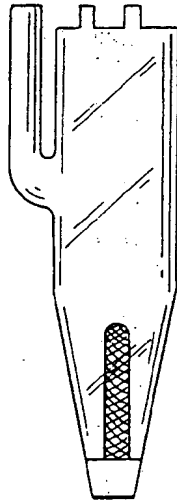


**Fig. 4**

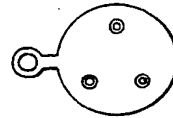




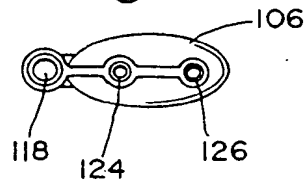
**Fig. 5**



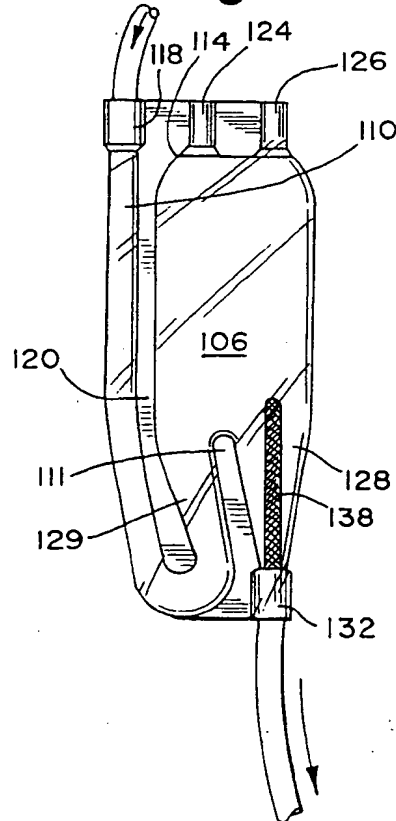
**Fig. 6**



**Fig. 9**



**Fig. 7**



**Fig. 8**

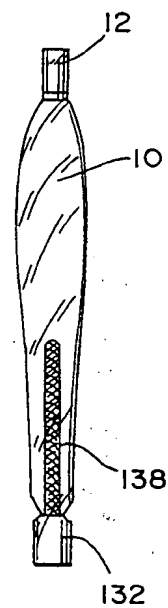


Fig. 12

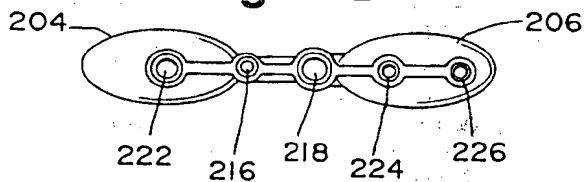


Fig. 10

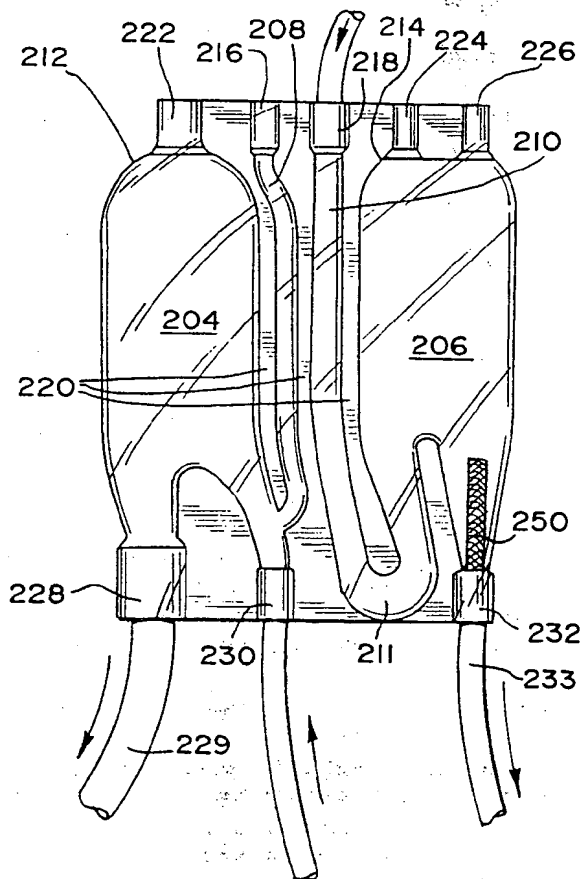


Fig. 11

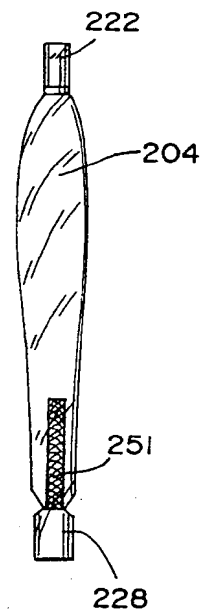
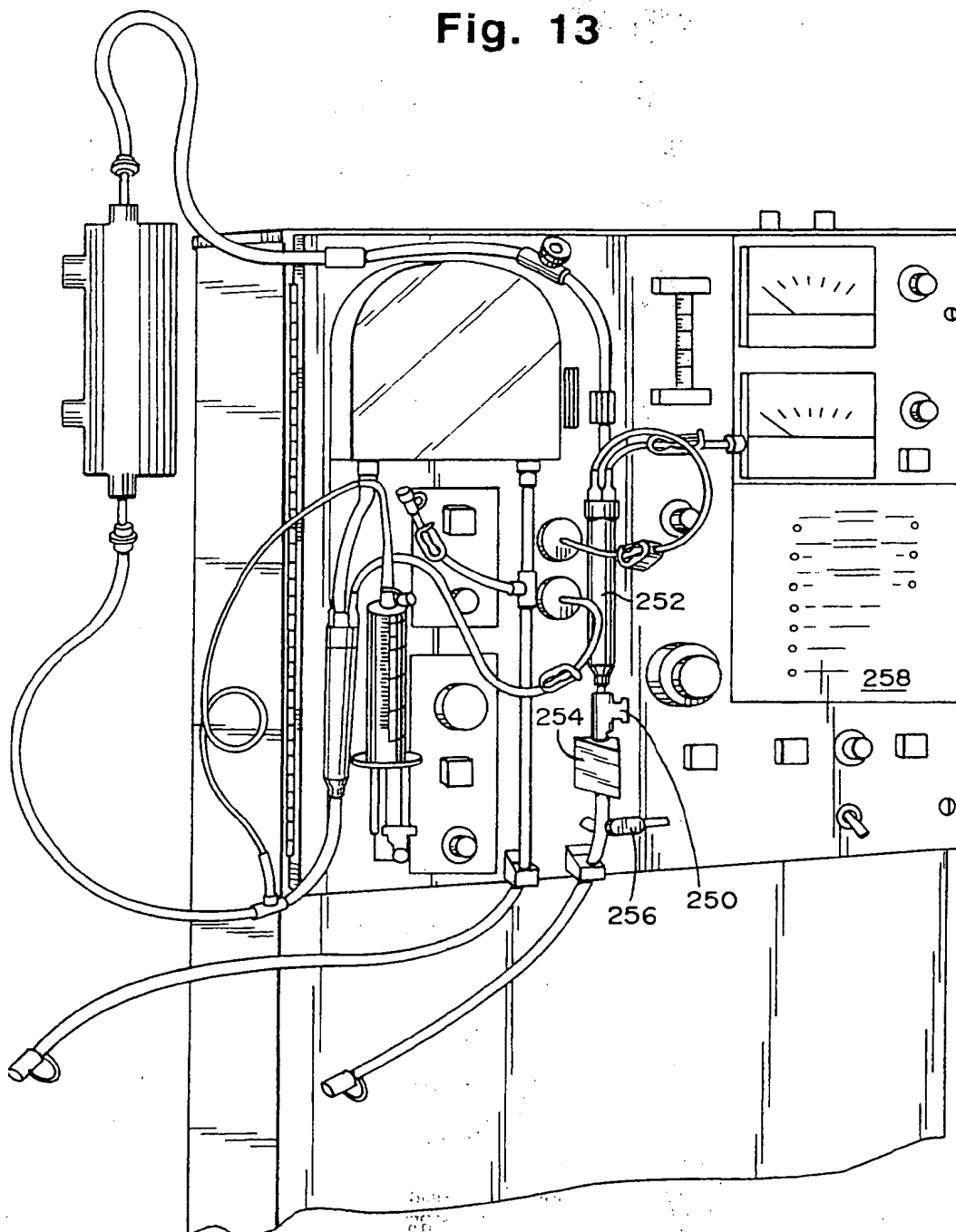


Fig. 13



1. The first part of the document discusses the importance of maintaining accurate records of all transactions. It emphasizes that proper record-keeping is essential for the transparency and accountability of the organization. This section also outlines the various methods used to collect and analyze data, ensuring that the information is reliable and up-to-date.

2. The second part of the document focuses on the implementation of the proposed changes. It details the steps involved in the transition process, from the initial planning phase to the final execution. This section also addresses the potential challenges and risks associated with the changes, providing strategies to mitigate them.

3. The third part of the document provides a comprehensive overview of the organization's current state. It includes a detailed analysis of the existing systems and processes, highlighting the strengths and weaknesses. This section also discusses the impact of the proposed changes on the organization's overall performance and the role of each department in the transition.

4. The fourth part of the document outlines the future goals and objectives of the organization. It describes the long-term vision and the specific targets that the organization aims to achieve. This section also discusses the resources required to achieve these goals and the timeline for the implementation of the changes.

5. The fifth part of the document provides a summary of the key findings and conclusions. It highlights the main points discussed in the document and provides a clear overview of the proposed changes. This section also includes a list of recommendations for the organization to follow, ensuring that the changes are implemented successfully.

6. The sixth part of the document provides a detailed description of the proposed changes. It includes a list of the specific changes to be made, along with the reasons for each change. This section also discusses the expected benefits of the changes and the potential risks associated with them.

7. The seventh part of the document provides a detailed description of the proposed changes. It includes a list of the specific changes to be made, along with the reasons for each change. This section also discusses the expected benefits of the changes and the potential risks associated with them.

8. The eighth part of the document provides a detailed description of the proposed changes. It includes a list of the specific changes to be made, along with the reasons for each change. This section also discusses the expected benefits of the changes and the potential risks associated with them.

9. The ninth part of the document provides a detailed description of the proposed changes. It includes a list of the specific changes to be made, along with the reasons for each change. This section also discusses the expected benefits of the changes and the potential risks associated with them.

10. The tenth part of the document provides a detailed description of the proposed changes. It includes a list of the specific changes to be made, along with the reasons for each change. This section also discusses the expected benefits of the changes and the potential risks associated with them.



(11) Publication number: **0 568 265 A3**

(12) **EUROPEAN PATENT APPLICATION**

(21) Application number: **93303126.2**

(51) Int. Cl.<sup>5</sup>: **A61M 1/36, A61M 5/165**

(22) Date of filing: **21.04.93**

(30) Priority: **30.04.92 US 876041**

(43) Date of publication of application:  
**03.11.93 Bulletin 93/44**

(84) Designated Contracting States:  
**AT BE CH DE DK ES FR GB IT LI NL PT SE**

(88) Date of deferred publication of search report:  
**27.09.95 Bulletin 95/39**

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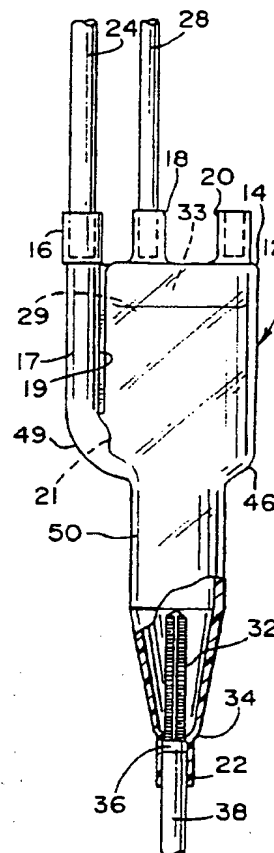
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(54) **Blow molded venous drip chamber for hemodialysis.**

(57) A blow molded, rigid plastic venous chamber (12) for hemodialysis, for use in a venous hemodialysis set (10). The chamber defines a plurality of first access ports (16,18,20) adjacent to a first end (14) thereof. A second access port (22) is provided at an opposed end. A tubular plastic filter (32) is positioned in close-fitting relation within the second access port to project into the chamber. Unique advantages are provided by such a blow-molded, filter-carrying chamber.

**Fig. 3**



**EP 0 568 265 A3**



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# EUROPEAN SEARCH REPORT

Application Number  
EP 93 30 3126

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.5)
A	GB-A-2 176 717 (COBE LABORATORIES, INC.) * page 2, line 22 - line 33; claims 1,3-5; figures 1-3 *	1,4-9, 11-15	A61M1/36 A61M5/165
A	EP-A-0 033 080 (ASID BONZ & SOHN GMBH) * page 2, line 34 - page 3, line 5 * * page 4, line 9 - page 5, line 7; claims 1,4; figures 10A,B,11,12 *	1-3,9-16	
A	DE-A-29 13 301 (BAXTER TRAVENOL LABORATORIES, INC.) * page 18, paragraph 3 - page 20, paragraph 2; claims 1,3,5; figures 1,3,5 *	1,4,7,8, 10,11	
A	US-A-5 061 365 (UTTERBERG) * column 1, line 64 - column 2, line 19 * * column 4, line 22 - line 29; claims 1,2; figures 1,2 *	1,4,5,11	
A	GB-A-2 047 542 (BAXTER TRAVENOL LABORATORIES, INC) * abstract; figures 1,3 *	17	<div>TECHNICAL FIELDS SEARCHED (Int.Cl.5)</div> <div>A61M</div>
The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 3 August 1995	Examiner Michels, N
<div>CATEGORY OF CITED DOCUMENTS</div> <div> X : particularly relevant if taken alone  Y : particularly relevant if combined with another document of the same category  A : technological background  O : non-written disclosure  P : intermediate document  T : theory or principle underlying the invention  E : earlier patent document, but published on, or after the filing date  D : document cited in the application  L : document cited for other reasons  &amp; : member of the same patent family, corresponding document </div>			

EPO FORM 1503/03.82 (P01C01)



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### CLAIMS INCURRING FEES

The present European patent application comprised at the time of filing more than ten claims.

- ☐ All claims fees have been paid within the prescribed time limit. The present European search report has been drawn up for all claims.
- ☐ Only part of the claims fees have been paid within the prescribed time limit. The present European search report has been drawn up for the first ten claims and for those claims for which claims fees have been paid, namely claims:
- ☐ No claims fees have been paid within the prescribed time limit. The present European search report has been drawn up for the first ten claims.

### LACK OF UNITY OF INVENTION

The Search Division considers that the present European patent application does not comply with the requirement of unity of invention and relates to several inventions or groups of inventions, namely:

see sheet -B-

- ☒ All further search fees have been paid within the fixed time limit. The present European search report has been drawn up for all claims.
- ☐ Only part of the further search fees have been paid within the fixed time limit. The present European search report has been drawn up for those parts of the European patent application which relate to the inventions in respects of which search fees have been paid, namely claims:
- ☐ None of the further search fees has been paid within the fixed time limit. The present European search report has been drawn up for those parts of the European patent application which relate to the invention first mentioned in the claims, namely claims:



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### LACK OF UNITY OF INVENTION

The Search Division considers that the present European patent application does not comply with the requirement of unity of invention and relates to several inventions or groups of inventions, namely:

1. Claims 1-16: Fluid collection chamber for an extracorporeal circuit.
2. Claim 17: Injection site for an extracorporeal circuit.